## IN THE CLAIMS

- 1. 14. (Cancelled)
- 15. (Previously Presented) A compound of formula (lb):

wherein R<sup>a</sup> is selected from halogen, hydroxy, cyano, nitro, amino, hydroxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkenyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, hydroxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylamino, di- C<sub>1</sub>-C<sub>6</sub> alkylamino, aminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, and C<sub>1</sub>-C<sub>6</sub> alkyltio; R<sup>b</sup> is selected from hydrogen, halogen, hydroxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxy cyano, nitro, amino, hydroxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkyl, perfluoro C<sub>1</sub>-C<sub>6</sub> alkyl, perfluoro C<sub>1</sub>-C<sub>6</sub> alkyl, di-C<sub>1</sub>-C<sub>6</sub> alkylamino, aminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, alkylaminoC<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub>

- 16. 17. (Cancelled)
- 18. (Previously Presented) A compound of formula (1b) according to claim 15 which is selected from the group consisting of: (2S,4S)-4-(2,3-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; and (2S,4S)-4-(2,5-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.
- 19. (Cancelled)
- 20. (Previously Presented) A pharmaceutical composition comprising a compound of formula (1b) according to claim 15, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

- 21. (Previously Presented) A pharmaceutical composition comprising a compound of formula (1b) according to claim 15, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.
- 22. (Previously Presented) A pharmaceutical composition according to claim 21, wherein the other therapeutically active agent is a PDEV inhibitor selected from sildenafil, vardenafil, tadalafil, 1-(6-ethoxy-5-[3-ethyl-6,7-dihydro-2-(2-methoxyethyl)-7-oxo-2H-pyrazolo[4,3-d]pyrimidin-5-pyridylsulfonyl]-4-ethylpiperazine, 5-(5-acetyl-2-butoxy-3-pyridinyl)-3-ethyl-2-(1-ethyl-3-azetidinyl)-2,6-dihydro-7H-pyrazolo[4,3-d]pyrimidin-7-one and 5-[2-ethoxy-5-(4-ethylpiperazin-1-ylsulphonyl)pyridin-3-yl]-3-ethyl-2-[2-methoxyethyl]-2,6-dihydro-7H-pyrazolo[4,3-d]pyrimidin-7-one.
- 23. 26. (Cancelled)
- 27. (Previously Presented) The compound (2S,4S)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid, or a pharmaceutically acceptable salt thereof.
- 28. (Withdrawn) The compound (2S,4S)-4-(2,3-Diffluoro-benzyl)-pyrrolidine-2-carboxylic acid, a pharmaceutically acceptable salt thereof.
- 29. (Withdrawn) The compound (2S,4S)-4-(2,5-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid, or a pharmaceutically acceptable salt thereof.
- 30. (Previously Presented) The salt, (2S,4S)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid mono hydrochloride salt.